delete

Fig. 3 (cf. example 6): Representation of the connection between viscosity of the peptide preparation and the standing time after reconstitution (viscosity was determined using a falling sphere microviscometer).

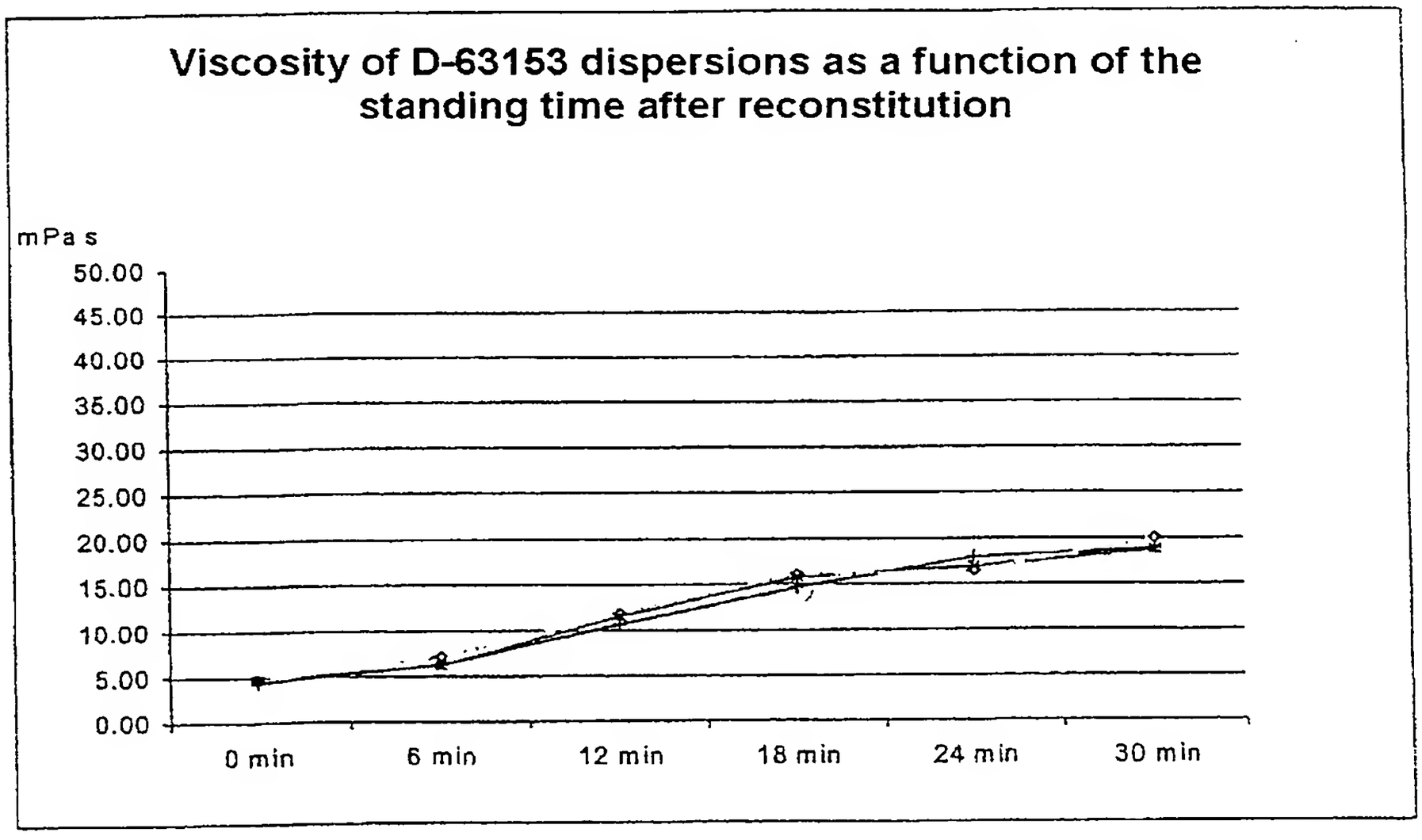

FIG. 3

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Fig. 4 (cf. example 7): Influence of the standing time after reconstitution on the plasma levels after s.c. injection; standing time = 0 min

Plasma concentration of D-63 153 after s.c. administration at 65 mg of D-63 153 dissolved in 2.6 nl of 0.1% (weight/ volume) NaCl solution; immediately after preparation of the test compound

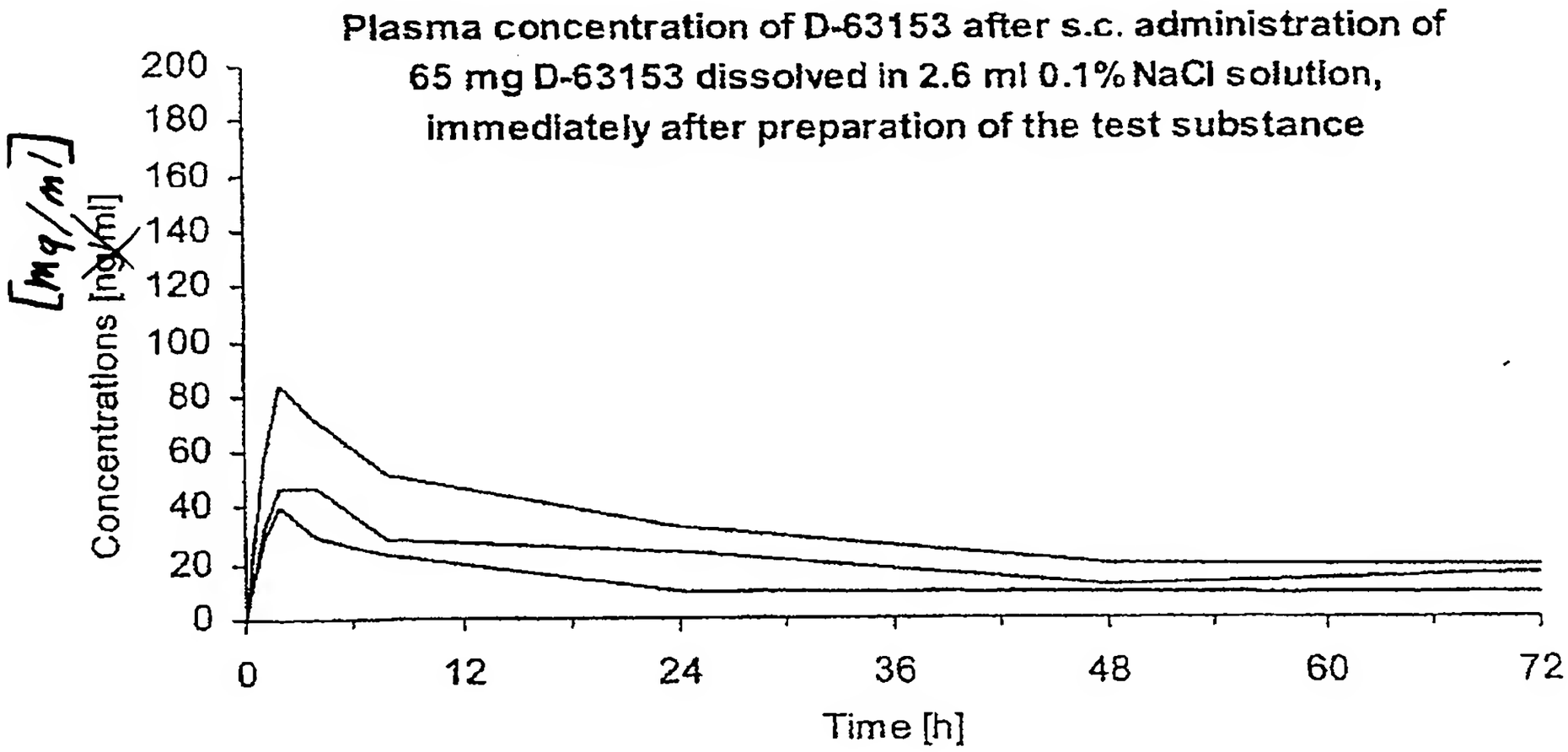

FIG 4

FIG. 4

Fig. 5 (cf. example 7): Influence of the standing time after reconstitution on the plasma levels after s.c. injection; standing time = 60 min.

Plasma concentration of D-63 153 after s.c. administration at 65 mg of D-63 153 dissolved in 2.6 nl of 0.1% (weight/volume) NaCl solution; 1 h (60 min) after preparation of the test compound

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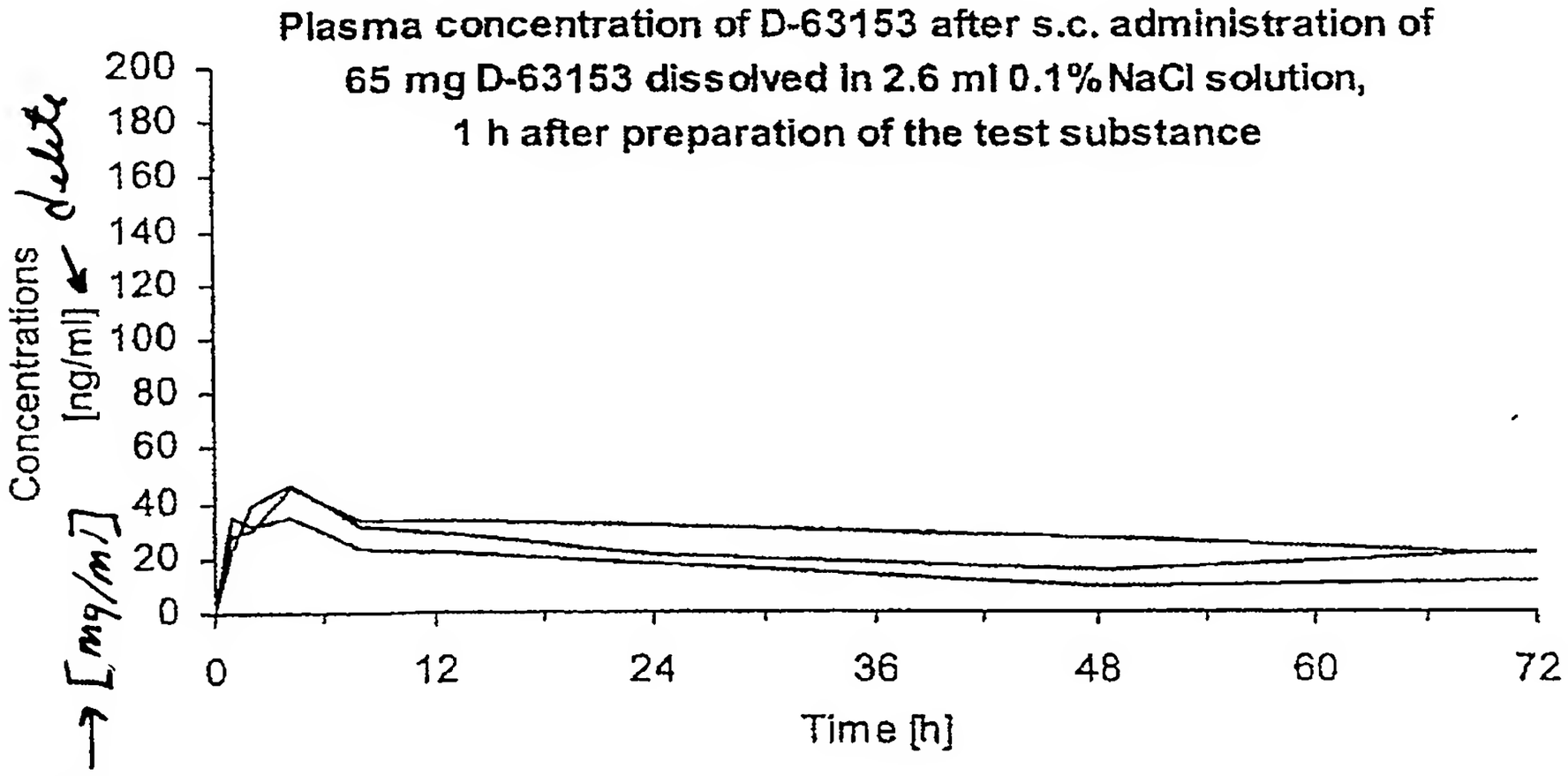

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FIG. 5